CLAIMS

We Claim:

- 1. A catalyst system comprising a cyclic germanium bridged bulky ligand metallocenetype catalyst compound and an activator.
- 2. The catalyst system of claim 1 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound has two bulky ligands.
- 3. The catalyst system of claim 2 wherein the bulky ligands are differently substituted.
- 4. The catalyst system of claim 1 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound is represented by the formula:

$$L^{A}(R'GeR')_{x}L^{B}MQ_{n}$$
 (I)

where M is a Group 3 to 7 transition metal, L^A and L^B is an unsubstituted or substituted, cyclopentadienyl ligand or cyclopentadienyl-type bulky ligand bonded to M; (R'GeR')_x is a cyclic bridging group bridging L^A and L^B, and the two R''s form a cyclic ring or ring system with Ge; independently, each Q is a monoanionic ligand, or optionally two Q's together form a divalent anionic chelating ligand; and where n is 0, 1 or 2 depending on the formal oxidation state of M, and x is an integer from 1 to 4.

- 5. The catalyst system of claim 4 wherein one of L^A or L^B is a substituted cyclopentadienyl or a substituted cyclopentadienyl-type bulky ligand.
- 6. The catalyst system of claim 1 wherein the catalyst system is supported.
- 7. The catalyst system of claim 4 wherein x is 1.
- 8. The catalyst system of claim 1 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound is represented by the formula:

 $-30 - L^{A}(R'GeR')_{x}L^{B}MQ_{n}$ (I)

where M is a Group 4, 5, 6 transition metal, L^A and L^B are bonded to M and are different, L^A and L^B are selected from the group consisting of unsubstituted or substituted, cyclopentadienyl ligands or unsubstituted or substituted, cyclopentadienyl-type bulky ligand; $(R'GeR')_x$ is a cyclic bridging group bridging L^A and L^B , and the two R''s form a cyclic ring or ring system with Ge; independently, each Q is a monoanionic ligand, or optionally two Q's together form a divalent anionic chelating ligand; and where n is 0, 1 or 2 depending on the formal oxidation state of M, and x is an integer from 1 to 4.

- 9. The catalyst system of claim 8 where x is 1.
- 10. The catalyst system of claim 8 wherein L^A and L^B are substituted or unsubstituted cyclopentadienyl rings.
- 11. The catalyst system of claim 8 wherein a least one of L^A and L^B is a cyclopentadienyl ring.
- 12. The catalyst system of claim 8 wherein L^A is a substituted cyclopentadienyl ring.
- 13. The catalyst system of claim 1 where the cyclic germanium bridged bulky ligand metallocene-type catalyst compound is selected from one of the group consisting of cyclotrimethylenegermyl(tetramethyl cyclopentadienyl) (cyclopentadienyl) zirconium dichloride, cyclotetramethylenegermyl (tetramethyl cyclopentadienyl) (cyclopentadienyl) zirconium dichloride, cyclotrimethylenegermyl(tetramethyl cyclopentadienyl) cyclopentadienyl) (2-methyl indenyl) zirconium dichloride, cyclotrimethylenegermyl(tetramethyl cyclopentadienyl) zirconium dichloride, cyclotrimethylenegermyl (tetramethyl cyclopentadienyl) (2,3,5-trimethyl cyclopentadienyl) zirconium dichloride, cyclotrimethylenegermyl bis(tetra methyl cyclopentadienyl) zirconium dichloride, cyclotetramethylenegermyl(tetramethyl cyclopentadienyl) zirconium dichloride, cyclotetramethylenegermyl bis(tetra methyl cyclopentadienyl) zirconium dichloride, cyclotetramethylenegermyl bis(tetra methyl cyclopentadienyl) zirconium dichloride, 3,4-dimethylcyclotetra-methyl-3-enegermyl(tetramethyl

cyclopentadienyl) (cyclopentadienyl) zirconium dichloride, 3,4-dimethylcyclotetramethyl-3-enegermylbis(tetramethyl cyclopentadienyl) zirconium dichloride, 3,4-dimethylcyclotetramethyl-3-enegermyl(tetramethyl cyclopentadienyl) (2,3,5-trimethyl cyclopentadienyl) zirconium dichloride, 3-methylcyclotetramethyl-3-enegermyl bis(tetra methyl cyclopentadienyl) zirconium dichloride, 3-methylcyclotetramethyl-3-enegermyl (tetra methyl cyclopentadienyl) (cyclopentadienyl) zirconium dichloride, 3-methylcyclotetramethyl-3-enegermyl (tetra methyl cyclopentadienyl) zirconium dichloride, o-xylidenegermyl bis(tetra methyl cyclopentadienyl) zirconium dichloride, o-xylidenegermyl(tetramethyl cyclopentadienyl) zirconium dichloride, and o-xylidenegermyl(tetramethyl cyclopentadienyl) (3-methylcyclopentadienyl) zirconium dichloride, and o-xylidenegermyl(tetramethyl cyclopentadienyl) (3-methylcyclopentadienyl) zirconium dichloride,

- 14. A process for polymerizing ethylene alone or in combination with one or more olefin(s) to produce a polymer product in the presence of a catalyst system comprising a cyclic germanium bridged bulky ligand metallocene-type catalyst compound and an activator.
- 15. The process of claim 14 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound has two bulky ligands.
- 16. The process of claim 14 wherein the bulky ligands are differently substituted.
- 17 The process of claim 14 wherein the polymer product is an ethylene copolymer having a I_{21}/I_2 greater than 35 and a melt strength greater than 7cN.
- The process of claim 14 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound is represented by the formula:

$$L^{A}(R'GeR')_{x}L^{B}MQ_{n}$$
 (I)

where M is a Group 3 to 12 transition metal, L^A and L^B is an unsubstituted or substituted, cyclopentadienyl ligand or cyclopentadienyl-type bulky ligand bonded to M; $(R'GeR')_x$ is a cyclic bridging group bridging L^A and L^B , and the two R''s form a

cyclic ring or ring system with Ge; independently, each Q is a monoanionic ligand, or optionally two Q's together form a divalent anionic chelating ligand; and where n is 0, 1 or 2 depending on the formal oxidation state of M, and x is an integer from 1 to 4.

- 19. A process for polymerizing ethylene alone or in combination with one or more other olefin(s) in the presence of a catalyst system comprising a cyclic germanium bridged bulky ligand metallocene-type catalyst compound and an activator, the process producing a polymer product having a melt strength greater than 7 cN and an I₂₁/I₂ of greater than 35.
- 20. The process of claim 19 wherein the catalyst system further comprises a carrier.
- 21. The process of claim 19 wherein the polymer product has a M_z/M_w greater than 3, and an I_{21}/I_2 of greater than 65.
- 22. A continuous gas phase process for polymerizing ethylene and at least one alpha-olefin having from 3 to 20 carbon atoms in the presence of a catalyst system comprising a cyclic germanium bridged bulky ligand metallocene-type catalyst compound, an activator and a carrier, the process producing a polymer product having a density greater than 0.900 g/cc, I₂₁/I₂ greater than 35, and a melt strength of greater than 6 cN.
- 23. The process of claim 22 wherein the I_{21}/I_2 is greater than 50.
- 24. The process of claim 22 wherein the cyclic germanium bridged bulky ligand metallocene catalyst compound is represented by the formula:

$$L^{A}(R'GeR')_{x}L^{B}MQ_{n}$$
 (I)

where M is a Group 3 to 7 transition metal, L^A and L^B are bonded to M and are different, L^A and L^B are selected from the group consisting of unsubstituted or substituted, cyclopentadienyl ligands or unsubstituted or substituted, cyclopentadienyl-type bulky ligand; $(R'GeR')_x$ is a cyclic bridging group bridging L^A and L^B , and the two R''s form a cyclic ring or ring system with Ge; independently, each Q is a monoanionic ligand, or optionally two Q's together form a divalent anionic chelating

ligand; and where n is 0, 1 or 2 depending on the formal oxidation state of M, and x is an integer from 1 to 4.

- 25. The process of claim 14 wherein the activator is alumoxane.
- 26. The process of claim 14 wherein the process is a gas phase polymerization process.
- 27. The process of claim 18 wherein M is a group 4, 5 or 6 metal.
- 28. A continuous gas phase process for the polymerization of olefin(s) to produce a polymer product in the presence of a catalyst system comprising a cyclic germanium bridged bulky ligand metallocene-type catalyst compound and activator.
- 29. The process in accordance with claim 34 wherein the activator is alumoxane.
- 30. The gas phase process in accordance with claim 34 wherein the olefin(s) comprises ethylene alone or in combination with one or more other olefin(s).
- 31. A process for polymerizing olefin(s) to produce a polymer product in the presence of a catalyst system comprising a saturatewd or monounsaturated cyclic germanium bridged bulky ligand metallocene-type catalyst compound and an activator.
- 32. The process in accordance with claim 37 wherein the activator is alumoxane.
- 33. The process of claim 38 wherein the cyclic germanium bridged bulky ligand metallocene-type catalyst compound is represented by the formula:

 $L^{A}(R'GeR')_{x}L^{B}MQ_{n}$ (I)

where M is a Group 3 to 12 transition metal, L^A and L^B is an unsubstituted or substituted, cyclopentadienyl ligand or cyclopentadienyl-type bulky ligand bonded to M; (R'GeR')_x is a cyclic bridging group bridging L^A and L^B , and the twoR''s form a saturated or monounsaturated ring or ring system with Ge; independently, each Q is a monoanionic ligand, or optionally two Q's together form a divalent anionic chelating ligand; and where n is 0, 1 or 2 depending on the formal oxidation statew of M, and x is an integer from 1 to 4.

- 34. An ethylene copolymer having a melt strength greater than 7cN and a I_{21}/I_2 greater than 35.
- 35. The ethylene copolymer of claim 25 wherein the I_{21}/I_2 is greater than 40.
- 36. The ethylene copolymer of claim 25 wherein the I_{21}/I_2 is greater than 60.
- 37. The ethylene copolymer of claim 25 wherein the I_2 is about 0.5 to 2 dg/min.
- 38. The ethylene copolymer of claim 25 wherein the ethylene copolymer has a density greater than 0.910 g/cc.
- 39. The ethylene copolymer of claim 25 wherein the ethylene copolymer is an ethylene/hexene-1 copolymer.